## AMENDMENTS TO THE CLAIMS

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- 1. (Original) A process for preparing optically active aldehydes or ketones which have from 3 to 25 carbon atoms and at least one racemizable stereocenter by catalytic dehydrogenation of the corresponding optically active primary or secondary alcohols in the gas phase in the presence of a catalyst.
- 2. (Original) The process according to claim 1, wherein a catalyst comprising at least one element selected from the group consisting of the elements calcium, zinc and copper is used.
- 3. (Currently amended) The process according to claim 1 [or 2], wherein the catalyst a eatalyst comprising zinc oxide and calcium carbonate is used.
- 4. (Currently amended) The process according to <u>claim 1</u>, <u>wherein the catalyst any of elaims 1 to 3</u>, <u>wherein a catalyst</u> whose active component comprises from 30 to 60% by weight of zinc oxide and from 40 to 70% by weight of calcium carbonates is used.
- 5. (Currently amended) The process according to claim 3 [or-4], wherein the calcium carbonate is present in the calcite modification.
- 6. (Currently amended) The process according to <u>claim 1</u> any of claims 1 to 5 for preparing branched or unbranched open-chain or monocyclic aldehydes or ketones.
- 7. (Currently amended) The process according to any of claims 1 to 6 claim 1 for preparing aldehydes or ketones which have a stereocenter in the  $\alpha$  and/or  $\beta$  position relative to the carbonyl group.
- 8. (Currently amended) The process according to any of claims 1 to 7 claim 1 for preparing optically active 2-methylbutan-1-al, 3,7-dimethyloct-6-en-1-al, 3,7-dimethyloctan-1-al, 8-p-menthen-3-one, p-menthan-3-one, 2-methylcyclohexanone, 3-methylcyclohexanone, 2-methylcyclopentanone, 3-methylcyclopentanone or 2,3-dimethylcyclohexanone.

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9. (Currently amended) The process according to any of claims 1 to 8 claim 1 for preparing optically active citronellal from optically active citronellal.

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- 10. (Currently amended) The process according to any-of claims 1 to 9 claim 1, wherein the enantiomeric excess (ee) of the aldehyde or ketone obtained corresponds to at least 90% of the enantiomeric excess of the alcohol used.
- 11. (Currently amended) The process according to any of claims 1-to 10 claim 1, wherein the dehydrogenation is carried out at a temperature in the range from 250 to 600°C.
- 12. (Currently amended) A process for preparing optically active menthol by cyclization of citronellal prepared according to any of claims 1 to 11 claim 1 to form isopulegol and subsequent hydrogenation.
- 13. cancelled

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- 14. (New) The process according to claim 2, wherein the catalyst comprising zinc oxide and calcium carbonate is used.
- 15. (New) The process according to claim 14, wherein the catalyst-whose active component comprises from 30 to 60% by weight of zinc oxide and from 40 to 70% by weight of calcium carbonates is used.
- 16. (New) The process according to claim 15, wherein the calcium carbonate is present in the calcite modification.
- 17. (New) The process according to claim 16 for preparing branched or unbranched openchain or monocyclic aldehydes or ketones.
- 18. (New) The process according to claim 17 for preparing aldehydes or ketones which have a stereocenter in the  $\alpha$  and/or  $\beta$  position relative to the carbonyl group.
- 19. (New) The process according to claim 18 for preparing optically active 2-methylbutan-1-al, 3,7-dimethyloct-6-en-1-al, 3,7-dimethyloctan-1-al, 8-p-menthen-3-one, p-menthan-3-one,

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2-methylcyclohexanone, 3-methylcyclohexanone, 2-methylcyclopentanone, 3-

methylcyclopentanone, 2,6-dimethylcyclohexanone or 2,3-dimethylcyclohexanone.

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